



Mountain-Pacific Quality Health Foundation

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*"The best quality
health care is provided to
every patient we serve,
every time."*

Montana Medicaid Prior Authorization Request Form for Use of Suboxone® (buprenorphine/naloxone) or Subutex® (buprenorphine) Coverage Restricted Exclusively for the Treatment of Opioid Addiction

Providers must submit their **Treatment Plan** before authorization will be considered.

At a minimum the **Treatment Plan** must include the following information:

- Documentation of assessment and screening for opioid dependence (DAST-10, DSM-IV)
- Documentation of opioid substance of abuse
- Documentation of proposed counseling schedule
- Documentation of proposed monitoring plan (urine drug screens, random pill counts, etc.)
- Copy of Controlled Substance/Treatment Contract which must include consequences for failure to comply

Patient's Name: _____ Date: _____

Patient I. D. Number: _____ D.O.B: _____

Physician's Name : _____

Physician's Phone # _____ Physician's Fax Number: _____

Drug/Dose Request: _____ (mg) Daily Directions: _____ (i.e.: 1 QD)

Is patient pregnant or nursing? _____ If Yes, Due Date? _____

Signature of Physician: _____

DEA#: X-_____ (Prescriber must have an X-DEA Number)

**Please complete form, attach documentation and fax to:
Medicaid Drug Prior Authorization Unit
1-800-294-1350**

Important Notice

The attached information is CONFIDENTIAL and is intended only for the use of the addressee(s) identified above. If the reader of this message is not the intended recipient(s) or the employee or agency responsible for delivering the message to the intended recipient(s), please note that any dissemination, distribution or copying of the communication is strictly prohibited. Anyone who receives this in error should notify us immediately by telephone, toll-free at (800) 395-7961 or locally at 406-443-6002 and return the original message to us at the address above via U. S. Mail.

Montana Medicaid Suboxone®/Subutex® Authorization Limitations

Covered Condition – Treatment of Opioid Addiction

Subutex®: Approvals will be limited to 5 days to allow for induction in the absence of a pregnancy diagnosis. For pregnancy, Subutex® will be authorized only for the duration of pregnancy. Maximum dose limitations for Suboxone® will apply.

Suboxone®:

- Patient must be 16 years or older.
- **Initial approval** –Prior authorization will be granted for 6 months. Dosing will be limited to maximum buprenorphine 24 mg/day. Requests for >24 mg/day will require provider documentation.
- *Documentation of compliance with all counseling, drug screen, and office visits must be provided for continuation of therapy beyond initial 6 months.*
- **Subsequent approvals-** Prior authorization will be granted for additional 6 month intervals up to 18 months to allow for a total of 24 months of therapy. Dosing will be limited to maximum buprenorphine 16 mg/day.
- Requests for dose increases will require provider documentation.
- Concurrent opioids, tramadol, or carisoprodol will not be covered. If a patient is Prior Authorized for Suboxone®/Subutex® after meeting all criteria and subsequently discontinues the medication, all opioids, tramadol formulations, and carisoprodol will remain on not-covered status. These medications will require Prior Authorization for any future prescriptions. Approval may be granted short-term for an acute injury, hospitalization, or other appropriate diagnosis only after the case is reviewed with the treating physician and the physician prescribing Suboxone®/Subutex®.

- **Note:** Approval may be cancelled at any time if patient fails to comply with **Treatment Plan: failure to attend counseling sessions; missed or inappropriate results from drug screens; breaking controlled substance/treatment contract.**